



S/N 10/044,796

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

|             |   |                 |             |
|-------------|---|-----------------|-------------|
| Applicant:  | LOSKUTOFF ET AL.  | Examiner:       | V. AFREMOVA |
| Serial No.: | 10/044,796  | Group Art Unit: | 1651        |
| Filed:      | OCTOBER 11, 2002  | Docket No.:     | 13511.1USU1 |
| Title:      | SEMEN EXTENDER COMPOSITION AND METHODS FOR<br>MANUFACTURING AND USING |                 |             |

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Declaration under 37 C.F.R. §1.131

I, Richard B. Lomneth, Ph.D. declare as follows:

1. I am one of the originally named inventors of the above-identified patent application.
2. Attached as Exhibit A are pages 3-6 from a laboratory notebook. These pages have been redacted to cover the dates in the upper right hand corners. The dates are all prior to May 14, 1999. The handwriting on the pages of laboratory notebook in Exhibit A is mine, and the reported compositions were prepared by me prior to May 14, 1999.
3. Sample 2 reported on laboratory notebook page 6 provided as Exhibit A identifies a sample I prepared containing 1 wt.% lecithin, 0.5 ml Biladyl® concentrate, 0.1 wt.% Equex, and water to 5 ml. To the 5 ml composition, 0.7 ml glycerol was added. Biladyl® is available from Minitüb GmbH, Germany. Biladyl® contains carbohydrate and buffer. Attached as Exhibit B is a product sheet for Biladyl®. In addition, Biladyl® is identified by the above-identified patent application at, for example, page 14, lines 15-18. Equex contains sodium lauryl sulfate as a surfactant and is identified by the above-identified patent application at page 16, line 11 through page 17, line 8 wherein "EQ" refers to Equex.

4. Sample 2 reported on laboratory notebook page 6 in Exhibit A describes a composition containing the components of independent claims 1 and 21 of the above-identified patent application. The phospholipid obtained from a non-animal source is satisfied by the lecithin, the surfactant to reduce ice crystal formation during freezing of the composition is satisfied by the sodium lauryl sulfate, the carbohydrate and the biological buffer are satisfied by the Biladyl® component, and the freeze agent is satisfied by glycerol. It is my belief that the composition of sample 2 exhibits a pH of about 6.9 to about 7.5 and an osmolality of about 250 mOsM to about 350 mOsM.

5. I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Date: \_\_\_\_\_

\_\_\_\_\_  
Richard B. Lomneth, Ph.D.

# START MAKING SAMPLES FOR TESTING

12-22 Bil A + 5% LECITHIN "BERRY BUFFAL BUFFAL"  
 12-23 " 3.5% " " " "  
 12-23 " 1% " " " "

- LECITHIN DISPERSED IN  $H_2O$ , THEN DILUTED INTO BILADYL A

BILADYL A - "10x" SOLUTION (40mL) DILUTE 400mL

SO LECITHIN PROBABLY DISPERSED IN  $H_2O$  TO H- FOR DILUTION OF BIL A CONCATIONS  
 BIL A + ADDITIVES MIXED 1:1 WITH SPERM + COOLED  
 BIL B = BIL A, <sup>TRAC</sup> DILUTED WITH GLYCEROL (8% LIGAND). MIX 1:1 WITH  
 SPERM / BIL A MIXTURE

EQUER PAPER "9-98"

CONCENTRATION = ? CALL MINI TUBE

DILUTE SERUM IN HEPES TL  
 ↳

NEED A BILADYL A STOCK

GENERAL PLAN:

1. MAKE A HEPES TL ANALOG FOR DILUTION OF SERUM SO ALWAYS SAME LECITHIN: SERUM DILUTION
2. MAKE LECITHIN, BIL A,  $H_2O$  AT HIGHER THAN FINAL CONCENTRATION; DILUTE IN BIL A 300mM AS NEEDED
3. MAKE LECITHIN-GLYCEROL STOCK SO THAT LECITHIN IS AT CORRECT FINAL CONCENTRATION (14% GLYCEROL RECOMMENDED BY DEVIN VOLLMER)
4. MAKE SAMPLES AT "PSEUDO CONCENTRATIONS" FOR THIS WEEK.

Paul & G

## 1. MAKE 0.1% EQUER PASTE IN LECITHIN SOLUTION

a. MAKE 10mL 1% LECITHIN, 0.1% EQUER

b. ~~2mL 5% LECITHIN~~

b. 10mL 1% LECITHIN

$$0.1\% \text{ EQUER PASTE} : C_1 V_1 = C_2 V_2$$

$$(10 \text{ mL})(0.1\%) = (100\%)(V_2)$$

$$10\% \text{ EQUER}$$

c. LABEL "1% LECITHIN, 0.1% EQUER"

2. CHOLESTEROL / METHYL- $\beta$ -CYCLODEXTRINa. USE ~5mg/mL Ch-Methyl  $\beta$  CYCLODEXTRIN WITH SAMPLEb. FOR 5mL SAMPLE, NEED 25mg CYCLODEXTRIN (= METHYL- $\beta$ -CYCLODEXTRIN)

c. STOCKS - 30mg CHOLESTEROL / 1% CYCLODEXTRIN = MBCO

d. MAKE 10mL IN  $H_2O$ 

$$(5\% \text{ CYCLODEXTRIN})(10 \text{ mL}) = .5\% \text{ CYCLODEXTRIN}$$

ADD ~15mg CHOLESTEROL

ADD  $H_2O$  TO 15mL TUBE, ADD 0.50 g METHYLATED  $\beta$  CYCLODEXTRIN  
CHOLESTEROL LOT J-8044

(- GOES INTO SOLUTION EASILY)

e. CHOLESTEROL: SUMMA C-3045 LOT 84H84585

43.8mg CHOLESTEROL

ADD 973  $\mu$ L ISOPROPANOL (+/- 5mg/mL IF DISSOLVED)

DISSOLVED AT ROOM TEMP W/ VORTEXING

f. IF 30mg CHOLESTEROL: 6.67mL

a. TUBE 10.67mL HAS 500mg CYCLODEXTRIN + 30mg CHOLESTEROL

WE WANT 5mg/mL

WANT 25mg CYCLODEXTRIN / 500  $\mu$ L = 46.86mg/mL

$$\left( \frac{46.86 \text{ mg}}{\text{mL}} \right) = \frac{25.0 \text{ mg}}{x} \quad x = 0.534 \text{ mL} / \text{TUBE} = 22 \text{ TUBES}$$

g. ADD CHOLESTEROL TO MBCO

- SLOW PROCESS - FIND WAY TO IMPROVE TEMP CONTROL MIXING

- SHOULD HAVE ADDED 15mg CHOLESTEROL !!!

- FIRST 15mg WENT IN EASILY; ALL BUT 0.080mL ADDED

- SPIN SAMPLE TO REMOVE UNDISSOLVED CHOLESTEROL. SPEC ALLOWED 0.080mL / TUBE

NOT EVERYTHING DISSOLVED - SPEED VAC / FINAL VOL = 1mL 350  $\mu$ L / TUBE

ALL WENT INTO SOLUTION

# TEST SOLUBILITY OF CHMBCA IN $H_2O$ , THEN DILUENT

- APPEARS TO COMPLETELY DISSOLVE IN 1.0 mL  $H_2O$ .
- ADD TO LECITHIN/BILANOL
  - APPEARS TO COMPLETELY DISSOLVE IN 1.0 mL 10% LECITHIN IN BILANOL

<sup>polyglycerol ester</sup>

- MAKE 5% PGE (SOLID, RAL) IN  $H_2O$   
 5g/100mL DO 2.1g/50mL OR 2.0g/40mL  
 (2.02g) ADD 40.0mL  $H_2O$

HEAT IN A MICROWAVE - OVERFLOWED - MICROSCOPICALLY  
 LOOKED WELL DISPERSED. STIRRING HIGH TEMP (NOT MEASURED TEMP)  
 - BETTER DISPERSED.

- MAKE 5% <sup>PGE</sup> IN BILANOL A (1/29 w/ ANTIBIOTICS)  
 2.1g PGE (SOLID) RL / 25mL BILANOL A

HEAT (IN MICROWAVE), STIRR ~ 2 MIN, SETTING 8  
 ↳ HOT (NOT MEASURED) TOO HOT TO HOLD COMFORTABLY

# TENTATIVE PROTOCOL FOR WORKING WITH SAMPLES.

1. MAKE SURE SERUM HAS  $225 \times 10^6$  MOTILE SPERM;  $>50\%$  MOTILITY
2. MIX SERUM WITH EQUINER, 1:1 RATIO
3. ~~COOL~~
4. MIX W/ 2ND SET OF EQUINER CONTAINING 14% GLYCEROL
3. FREEZE

$$\frac{14\% \times 5\%}{5\%} = 14$$

$$\frac{14\% \times 5\%}{5\%} = 14$$

$$\frac{14\% \times 5\%}{5\%} = 14$$

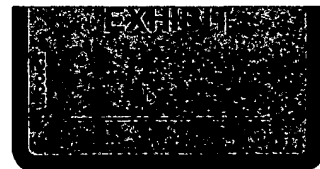
$$\frac{14\% \times 5\%}{5\%} = 14$$

ADD 0.80mL  
GLYCEROL

PGE APPEARS TO  
HAVE PRECIPITATED OUT OF THEM  
SOLUTIONS MAY HAVE TO BE  
A LOWER MW (MORE SOLUBLE) FORM

## SAMPLES

- ✓ 1. 5mL 1% LECITHIN/BILADYL + 0.7mL GLYCEROL (5mL 1% Lec/Bil From 12.23)
- ✓ 2. 5mL 1% LECITHIN/BILADYL + 0.1% EQUINER + 0.7mL GLYCEROL (5mL 1% Lec/EQUINER/Bil From 3.15.99)
- ✓ 3. 5mL 1% Lec/Bil + 0.7mL GLYCEROL - ADD 1mL TO CH-MPCD TUBE & DISPERSE  
MIX WITH REMAINDER OF 1% Lec BILADYL (5mL 1% Lec/Bil From 12.23)
- ✓ 4. 5mL 1% Lec/Bil + 0.1% EQUINER + 0.7mL GLYCEROL - ADD 1mL TO CH-MPCD TUBE, DISPERSE  
+ MIX W/ REMAINDER (5mL 1% Lec/EQUINER/Bil From 3.15.99)
- ✓ 5. SAME AS "3" EXCEPT USE 2 TUBES CH-MPCD (1% Lec/Bil From 12.23)
- ✓ 6. 5mL 4% PGE, 1% Lec/Bil + 0.7mL GLYCEROL (4mL 5% PGE/Bil + 1mL 5% Lec 12.22)
- ✓ 7. 5mL 2% PGE, 1% Lec/Bil + 0.7mL GLYCEROL (2mL 5% PGE/Bil + 1mL 5% Lec 12.22 + 2mL Bil 12.24)
- ✓ 8. 5mL 4% PGE, 1% Lec/Bil + 0.1% EQUINER + 0.7mL GLYCEROL (SAME AS 6 PLUS 5mL EQUINER)
- ✓ 9. 5mL 2% PGE, 1% Lec/Bil + 0.1% EQUINER + 0.7mL GLYCEROL (SAME AS 7 PLUS 5mL EQUINER)
- ✓ 10. 5mL 4% PGE, 1% Lec/Bil + ADD TO CH-MPCD TUBE AS ABOVE (SAME AS 6 PLUS CH-MPCD)
- ✓ 11. 5mL 2% PGE, 1% Lec/Bil + ADD TO CH-MPCD TUBE AS ABOVE (SAME AS 7 PLUS CH-MPCD)
- ✓ 12. 5mL BILADYL NO LECITHIN + 0.7mL GLYCEROL (5mL BILADYL w/ ANTIBIOTICS 1.24)
- ✓ 13. 5mL 4% PGE, 1% Lec/Bil, 0.1% EQUINER, ADD TO CH-MPCD AS ABOVE (SAME AS 8)
- ✓ 14. 5mL 2% PGE, 1% Lec/Bil, 0.1% EQUINER, ADD TO CH-MPCD AS ABOVE (SAME AS 9 PLUS CH-MPCD)



## Preparation of COCKTAIL AB:

Add 12 ml of double distilled, sterile water, using a sterile syringe.

**Final composition of reconstituted COCKTAIL AB expressed as active units of antibiotics per 0.02 ml:**

100 µg Tylosin,  
500 µg Gentamicin,  
300 µg Lincomycin,  
600 µg Spectinomycin

## Usage for „Neat Semen Treatment“:

Add and carefully mix 0,02 ml to each ml of neat semen, using a sterile syringe.

## Usage for BILADYL SOLUTION A:

Add and carefully mix 10 ml to SOLUTION A, using a sterile syringe.

## Preparation of SOLUTION A:

- 1) Reconstitute 49 g of SOLUTION A with double distilled sterile water to a combined volume of 390 ml.
- 2) Add 100 ml clean yolk from fresh chicken eggs.
- 3) Add 10 ml of reconstituted antibiotics COCKTAIL AB, using a sterile syringe.
- 4) Mix gently and warm mixture to + 30° C (+ 86° F)
- 5) Filter medium before adding it to semen.

## Preparation of SOLUTION B:

- 1) Reconstitute 250 g of SOLUTION B with double distilled sterile water to a combined volume of 400 ml.
- 2) Add 100 ml clean yolk from fresh chicken eggs.
- 3) Mix gently and warm mixture to + 30° C (+ 86° F)
- 4) Filter medium before adding it to semen.

## Usage:

Dilute semen with equal quantities of SOLUTION A and B according to the CSS Processing Regulations.

## Final composition of SOLUTION A and B per 100 ml, as approved by CSS:

Yolk 20 %, Glycerol 7 %, Tris 2,42 %, Citric Acid 1,38 g%, Fructose 1,00 g%,  
Active Units of Antibiotics:  
Tylosin 5,25 mg, Gentamicin 26,25 mg, Lincomycin 15,75 mg, Spectinomycin 31,5 mg and double distilled sterile water

## Storage:

At a controlled temperature of + 5° C (+ 41° F) in a dark environment.  
Shelf life: 12 months.

## Warning:

Keep out of reach of children  
Not for human or animal consumption and/or treatment  
Not for injection  
Not for use on live animals  
Do not expose to heat or sun

Made in Germany

**BILADYL IS APPROVED BY CERTIFIED SEMEN SERVICES INC.**



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